

(12)

EUROPEAN PATENT APPLICATION

(21) Application number: 85108192.7

(51) Int. Cl.⁴: **A 61 K 7/20**
B 65 D 35/00

(22) Date of filing: 02.07.85

(30) Priority: 23.05.85 US 737157
17.06.85 US 745993

(43) Date of publication of application:
26.11.86 Bulletin 86/48

(84) Designated Contracting States:
DE FR GB IT

(71) Applicant: **Schaeffer, Hans A.**
14 Pallant Avenue
Linden New Jersey(US)

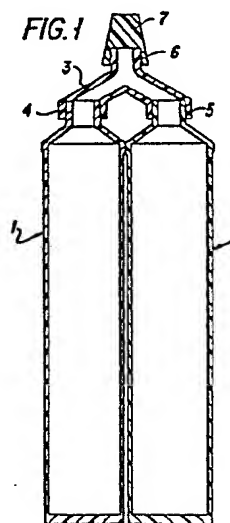
(72) Inventor: **Schaeffer, Hans A.**
14 Pallant Avenue
Linden New Jersey(US)

(74) Representative: **Patentanwälte Grünecker, Kinkeldey,**
Stockmair & Partner
Maximilianstrasse 58
D-8000 München 22(DE)

(54) **Dental preparation, article and method for storage and delivery thereof.**

(57) Disclosed is a composition useful in combatting gum disease including (a) a gel component having (i) about 0.1 - 10% by weight of hydrogen peroxide; (ii) about 0.05 - 5.0% by weight of a water-dispersible copolymer of acrylic acid cross-linked with polyallyl sucrose; (iii) about zero to about 2.0% by weight of a nonionic cellulose stabilizer; (iv) a neutralizing agent selected from the group consisting of sodium hydroxide, potassium hydroxide, triethanolamine, diisopropylamine and ammonia in an amount sufficient to raise the gel pH to about 3 - 6; and (v) purified water; and (b) a paste component having: (i) about 2 - 60% sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, MgCl₂, MgSO₄, and K₂SO₄; (iii) about 2 - 60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol and mixtures thereof; (iv) about 0.1 - 5% of a thickener stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium oxide and mixtures thereof; and (vi) purified water. The past component and gel component are combined immediately prior to use.

Also disclosed are combinations of a two part container and a composition of a gel component and a paste component.



1 European Patent Application No. 85 108 192.7
Applicant: Schaeffer, Hans A.

5

10

DENTAL PREPARATION, ARTICLE AND METHOD
FOR STORAGE AND DELIVERY THEREOF

15

This invention relates to a dental preparation useful in the treatment of gum disease, to a method of storing and delivering such preparation to a use point and to an article for the storage and delivery of such preparation.

20

25

It has long been recognized that the combination of hydrogen peroxide solution with sodium bicarbonate and table salt has an excellent curative and preventive effect on gum disease caused by bacterial infection. Dr. Paul H. Keyes has advocated use of this combination to the dental profession and to the public at large based on his work of more than 25 years on the subject, which has shown that upon daily and diligent topical application of these materials, gum disease may be effectively controlled. On the basis of his recommendations, many dentists urge their patients to use the Keyes procedure (substantially as described e.g. in S. Elder: "An Alternative To Gum Surgery" Modern Maturity, Aug.-Sept. 1980 pp. 31-32).

30

35

Dr. Keyes advocates that a quantity of solid sodium bicarbonate be placed in one hand, and that the toothbrush, held in the other hand, be dipped into a hydrogen peroxide - table salt solution and then transferred to the bicarbonate and applied to the teeth and gums. Upon contact with the gums, the hydrogen peroxide is exposed to the enzyme catalase, which is always present

1 in the buccal cavity, and is attacked thereby resulting
in the release of active oxygen. The combination of the
active oxygen and the sodium bicarbonate together with
table salt destroys the bacteria responsible for gum
5 disease. Unfortunately, hydrogen peroxide and sodium
bicarbonate may not be premixed, as they immediately react
and are thereby rendered ineffective against gum disease.
In addition, hydrogen peroxide is unstable and therefore
difficult to store for prolonged periods of time. Finally,
10 mere dipping of the toothbrush in a hydrogen peroxide
solution does not insure delivery of a sufficient amount
of hydrogen peroxide to the teeth and gums. These factors
are responsible for the fact that use of the Keyes
procedure is extremely awkward, inconvenient and messy.
15 Another disadvantage stems from the fact that, the mixture
of hydrogen peroxide and sodium bicarbonate has a very
unpleasant taste. For these reasons, patients have shown
extreme reluctance to follow this procedure, especially on
a daily basis, as would be required for effective gum
20 disease control. As a result, the benefits which the
Keyes procedure affords have largely been left unrealized.

Accordingly, it is an object of this invention to
eliminate the above disadvantages associated with use of
the Keyes procedure by providing a dental preparation
25 incorporating the active constituents of the Keyes pro-
cedure that has pleasant taste and is neat and convenient
to use, and a method for using such preparation that
permits contact between hydrogen peroxide and sodium
bicarbonate only shortly before use and, therefore,
30 assures maximum effectiveness against gum disease.

It is another object of the present invention
to provide a dental preparation incorporating the active
constituents of the Keyes method and a method for using
such preparation that permits a sufficient, consistent and
35 reproducible amount of hydrogen peroxide to be delivered to
the use point.

1 It is yet another object of this invention to
provide an article for the storage and delivery of this
improved dental preparation which makes its use neat and
convenient and which prevents contact between hydrogen
5 peroxide and sodium bicarbonate prior to application.

 In accordance with the present invention,
hydrogen or urea peroxide is dissolved in a nontoxic gel
for use in combination with a separately stored but
substantially simultaneously dispensed paste containing
10 sodium bicarbonate, table (or another suitable) salt, and,
preferably, additional cleansing, anticaries and polishing
agents as well as an effective concentration of flavoring
substances. Each of the gel and paste are loaded either
into separate collapsible containers which are connected
15 by means of a common orifice (as in Figure 1), or which
have substantially adjacent orifices (as in Fig. 2), or in
separate compartments of a single container (as in Figs. 3
or 4). Alternatively, the gel and paste may be loaded in
separate compartments of a two-compartment pressurized
20 container (as in Fig. 5) or a mechanically actuated pump,
as in Figure 6.

 Upon substantially simultaneous squeezing of the
containers, in much the same way as common toothpaste
tubes (or upon actuating of the pressurized container or
25 pump), controlled quantities of the gel and paste can be
simultaneously released onto the toothbrush and immedi-
ately applied to the teeth and gums. Control of the
peroxide, salt, and NaHCO_3 quantities delivered may be
thus effected by specification of the opening of the
30 orifice and the active ingredient concentration in each
tube (or pump compartment). As described above, when the
brush is applied to teeth and gums, immediate mixing of
the products takes place followed by the rapid evolution
of active oxygen and carbon dioxide. At the same time,

1 the effervescence accompanying release of active oxygen
activates the flavor contained in the bicarbonate paste
and produces a lasting highly refreshing taste in the
mouth which is unlike any other flavor provided by exist-
5 ing toothpastes or gels.

Another advantage afforded by the present
invention, as compared with the Keyes procedure, is that a
greater and more consistent amount of peroxide is delivered
to the use point.

10 Yet another advantage stems from the tendency of
the present composition to cling to the gum tissues and
thus provide them with the full benefit of substantially
all of the composition applied to the gums.

Gelling agents suitable for use in preparation
15 of the peroxide gel in accordance with this invention
should be nontoxic and neutral to the peroxide to assure
its stability. In addition, they should be preferably
sensitive to external electrolytes, such as those con-
tained in the sodium bicarbonate paste, in order to make
20 peroxide immediately available to the oral tissues. A
gelling agent suitable for use with the present invention
is a copolymer of acrylic acid cross-linked with polyallyl
sucrose, as described in U.S. Patent No. 2,798,053
issued on July 2, 1957 and assigned to B. F. Goodrich
25 Inc. Other gelling agents resulting in stable hydrogen
(or urea) peroxide gels suitable for use in the present
invention include those described in British Patent No.
827,331, i.e., organic polymeric acid colloids including
polyuronic acids, carboxypolymethylene compounds and
30 polyester resins containing three carboxyl groups, such as
partially hydrolized polyacrylates or polymethacrylates
and copolymers thereof; and those described in U.S. Patent
No. 3,639,574 issued on February 1, 1972 to Schmolka, i.e.,
polyoxyethylene polyoxypropylene block copolymers, which,

1 according to Schmolka, may be used in the preparation of
stable, firm hydrogen peroxide gels. Preferred are
water-dispersible copolymers of acrylic acid cross-linked
with about 0.75 to about 1.5 % of polyallyl sucrose and
5 neutralized with triethanolamine, NaOH or another alkaliz-
ing agent, as taught in U.S. Patent No. 3,499,844^{1/}
issued on March 10, 1970 to Kibbel et al. For purposes of
the present invention, Kibbel's acrylic copolymer may
be preferably combined with an anionic or non-ionic
10 surfactant, such as disclosed in U.S. Patent No. 4,130,501^{2/}
issued on December 19, 1978 to Lutz et al. Such surfac-
tants are not essential for the formation of a stable
hydrogen peroxide gel in accordance with this invention,
but may be added to facilitate distribution and rapid
15 penetration of the active oxygen throughout the area to be
treated. A particularly preferred gelling agent for the
purposes of the present invention is that described by
Kibbel, supra. This gelling agent may but does not have
to be modified by the addition of a suitable amount of
20 non-ionic cellulose gum such as hydroxyethyl- or hydroxy-
propyl-cellulose or hydroxypropyl-methyl-cellulose in
order to improve the physical stability of the gel,
especially when subjecting it to stress such as that
resulting from squeezing of the collapsible tubes, or
25 pumping action.

The most preferred gelling agents are marketed
under the trademark CARBOPOL 941 or 1342 by Goodrich.
Carbopol 941 does not need neutralization for gelling (and

30 _____
^{1/} The disclosures of these patents are incorporated
herein by reference.

^{2/} The disclosures of these patents are incorporated
35 herein by reference.

1 preferably is not neutralized in this invention) because
it gels readily in the presence of hydrogen donors.
Carbopol 941 has proved to have greater long term physical
stability (also believed to be due to hydrogen bonding).
5 Although Carbopol 1342 has just become available on the
market and its composition and characteristics have not
been fully disclosed, it is claimed by the manufacturer
that this acrylic acid copolymer (even though it needs to
be neutralized) displays satisfactory long term stability
10 comparable to that of Carbopol 941.

Gels made from these agents do not need any
cellulose additive as a stabilizer, because they are
thixotropic (and also pseudoplastic).

Not only is Carbopol 941 the most preferred
15 gelling agent for non-neutralized gels, it is also most
preferred for neutralized gels along with Carbopol 1934,
940 and 1342.

The hydrogen peroxide gel may then contain the
following ingredients in the following amounts - H_2O_2 :
20 about 0.1 - 10.0% and preferably about 3.0 - 6.5%; Acrylic
acid copolymer: about 0.05 - 5.0% and, preferably, about
1.0 - 3.0%; nonionic cellulose gum (optional): about 0 -
2.0% and, preferably, about 0.3 - 1.5%; neutralizing agent
(triethanolamine, diisopropanolamine, NaOH, KOH): an
25 amount sufficient to raise the gel pH to about 3.0 - 6.0;
NaOH is preferred. The balance is purified (distilled or
deionized) water.

If a non-neutralized gelling agent is used, the
aforementioned gel may contain about 2 - 80% and prefer-
ably about 20 - 60% by weight of a polyol selected from
30 the group consisting of glycerin, sorbitol (70% solution)
propylene glycol, polyethylene glycol, ethoxylated or
propoxylated lower ($C_2 - C_5$) fatty alcohols and
mixtures thereof. The preferred polyol is glycerin.
35 The amount of the water is decreased so that the total

-7-

1 adds up to 100% by weight. The pH need not be controlled
but falls between about 2 and 4.

5 The sodium bicarbonate paste contains sodium
bicarbonate, sodium chloride (or another suitable salt
although the salt may be omitted, if desired), purified
(distilled or deionized) water, a thickener/stabilizer
such as cellulose gum and or magnesium-aluminum silicate,
as essential ingredients and, most preferably, it also
contains a polishing/stabilizing agent, such as bentonite,
10 silica, titanium dioxide, magnesium oxide or mixtures
thereof (the first three and their mixtures are preferred).
In order to disperse the "chalky" taste imparted mostly by
the bicarbonate and enhance the taste and plasticity of
the paste, a bodying agent is added, such as sorbitol,
15 glycerin and/or a glycol. In addition, if the paste (in
combination with the gel) is to displace toothpaste
completely, additional cleansing agents, such as calcium
sulfate, calcium phosphate, hydrated aluminum oxide,
calcium carbonate, magnesium carbonate, and magnesium
20 silicate or mixtures thereof can be added. A fluorine-
containing compound is also preferably included for its
anti-carries activity. Suitable fluorine-containing
compounds are NaF, Na-monofluorophosphate, KF, potassium
monofluorophosphate, sodium fluorosilicate, sodium
25 fluorozirconate, etc. (with NaF being most preferred).
Finally, a foaming agent such as sodium lauryl sulfate
(most preferred), sodium N-lauroyl sarcosinate, sodium
coconut monoglyceride sulfonate, sodium N-methyl-N-palmi-
toyl lauride or a nonionic surfactant such as a polysor-
30 bate (e.g., Tween 60 or 80 manufactured by ICI Americas,
Wilmington, Delaware) or poloxamer or mixtures thereof,
which also enhances the peroxide-bicarbonate-salt action,
may be added. Flavoring agents, such as sodium saccharin,
or other artificial sweeteners, peppermint or spearmint or

1 other flavors are preferably added to further curb the
unpleasant taste. Finally, methyl, butyl and/or propyl
paraben, sodium benzoate, potassium sorbate or mixtures
thereof are preferably added as preservatives, with methyl
5 and propylparaben being most preferred. Use of a coloring
agent is optional.

The constituents and quantities for the bicarbonate paste are as follows:

sodium bicarbonate: about 2 - 60% and preferably
10 20 - 40%;

salt: about 0 - 6%, preferably about 1 - 6% and most
preferably about 2 - 4% of NaCl (preferred) or KCl,
MgCl₂, MgSO₄, Na₂SO₄ or K₂SO₄ or mixtures
thereof;

15 humectant: about 2 - 60% and preferably, 15 - 25%
consisting of glycerin, sorbitol, polyethylene glycol,
polypropylene glycol, ethoxylated or propoxylated lower
fatty alcohols and mixtures thereof;

thickener-stabilizer: nonionic cellulose gum -- about
20 0.1 - 5% and preferably 1.0 - 2.0%; or magnesium aluminum
silicate or mixtures thereof in the same proportions;

stabilizer/polishing agent/cleanser: total about 1 -
30%, preferably about 1.5 - 20%; these preferably include
one or more of: bentonite -- about 0.5-7.5%; silica --
25 about 0.1-8.0%; titanium dioxide -- about 0.1-8.0%; and/or
magnesium oxide -- about 0.2-8.0%; preferably, about
1.5-5.0; 0.5-6.0; 0.5-3.0; and 0.5-3.0 percent, respec-
tively.

fluorine-containing compound: sufficient to yield 200
30 to 3,000 ppm and preferably 1,000 to 2,000 ppm fluorine;

foaming agent: about 0.1 - 2.5%; preferably about
0.2 - 0.5%;

additional polishing agents: total about 1 - 30%,
preferably about 5 - 20%;

35 flavoring agent(s): to taste, preferably 0.1 - 2%;

1 preservatives: about 0.05 - 0.5%.

 The balance is purified water. A coloring agent may be added. The paste and the gel are preferably used in substantially equal proportions, by volume.

5 If urea peroxide is used in the gel, the bicarbonate paste composition does not change. The gel composition will be as follows:

urea peroxide: about 2 - 25%, preferably about 8 - 12%;

acrylic copolymer: about 0 - 3.5, preferably about 1 - 3%;

10 glycerin: balance.

 The other polyols described above are reactive with the urea peroxide and should not be used.

 The gel and paste combination may be simultaneously dispensed from separate collapsible tubes

15 preferably made of plastic, or a plastic/metal laminate (to avoid reaction with H_2O_2) and enhanced flavor retention), such as tubes 1 and 2 shown in Figure 1.

 The tubes are fitted with a Y-shaped conduit 3 which provides them with a common orifice 6. Conduit 3 may also
20 be made of plastic (preferably by injection molding) and is preferably detachably but snugly attached to mouths 4,5 of tubes 1,2 so that it may be removed for cleaning. For additional convenience and in order to ensure dispensation of substantially equal amounts of the gel and paste, the
25 tubes themselves may be held together, e. g., by banding or cementing, along corresponding dorsal sides, shown in Fig. 1, or, preferably, along corresponding ventral sides (see, e. g., Fig. 3A).

 Alternatively, the two tubes may be constructed
30 to have a common (preferably flat) sidewall portion 26 as shown in Figure 2. In the latter case, the Y-shaped conduit may be unnecessary, if the mouths 24,25 of tubes 21,22 are sufficiently close so that sufficient quantities of the gel and paste may be simultaneously dispensed
35 directly on the toothbrush. Conventional toothpaste or

-10-

1 medicament tubes may be thus used after one of their side
walls and the corresponding portion of their head struc-
ture are permanently deformed (e.g. by application of
pressure) to a substantially flat surface.

5 A third alternative packaging method involves
loading each of the gel and paste into separate compart-
ments of the same collapsible tube, joined by a common
orifice, as shown in Figure 3. Composite tube 31 has
compartments 32, 33 separated by divider 34 which is
10 firmly attached along substantially diametrically opposed
portions 35,36 of the sidewall 37, and corresponding
portions 38,39 of head structure 40. Divider 34 may be
glued or welded to sidewall 37 and head structure 40 of
tube 31 during manufacture of the latter. Divider 34 is
15 preferably provided with protruding portion 41, which
extends into the mouth 42 of tube 31 until its edge is
substantially flush with rim 43 of mouth 42. Thus,
divider 34 forms with sidewall 37 two separate compart-
ments 32,33 of substantially the same volume for storage
20 of the gel and paste, respectively.

In another alternative packaging method, the two
tubes are "concentric" as shown in figure 4A. Inner tube
401 lies within and parallel with outer tube 402.

25 The mouths of the tubes 401 and 402, designated
as 405 and 404 in Figure 4, abut at the same point. Tube
401 is fastened on to tube 402 at the mouth by protrusion
406 (shown in enlargement in Fig. 4B, which is a cross-
section of the embodiment of Fig. 4A taken along 4B-4B).
30 Protrusions 406 are inserted in a groove of mouth 404 of
tube 402. The material contained in tube 402 can still
pass through the available space between mouth 404 of
outer tube 402 and mouth 405 of inner tube 401. Engage-
ment of pins 406 in the groove secures the inner tube 401
35 on the outer tube 402.

-11-

1 The closure 407 of the tube-within-a-tube (which
can screw on the outer tube or simply be held by pressure)
arrangement may but does not have to be equipped with an
interior protrusion 408 to fit in the inner tube in order
5 to prevent premature intermixing of the two components at
the mouth of the tube. Because of the pseudoplastic
quality of the gel and/or the memory of the plastic
tube, however, such intermixing is not likely to occur.
The tubes are filled from the bottom and are (subse-
10 quently) sealed together by conventional techniques.

Other alternative packaging arrangements are
disclosed in Figures 5 and 6. Pressurized container 501
in Figure 5 is provided with two compartments 502 and 503
and two spouts 504 and 505. The internal pressure of the
15 compartments is maintained by pressurized nitrogen, at the
bottom 506 of each compartment. Operation of the mechanical
actuator 507 (by pressing downwards) actuates valves
508 and 509 which release the contents of the compartments
through the spouts (channels) causing the upwardly
20 slidable sealing disks 510 and 511 (guided by members 512,
513) to move up along the compartments (due to the nitrogen
being under pressure). Similar (but conventional)
pressurized containers are manufactured for example by
American Can Company. A dual compartment container, as
25 described above, would be a modification of the existing
containers.

In an alternative pump embodiment depicted in
Figs. 6 and 7 a pressurized container 601 is provided with
two compartments 602 and 603, and two spouts 604 and 605
30 for dispensing the gel and paste. Located within the tube
605 is a first piston 606 which is held in place by the
upper surface of the contents within compartment 603 and a
tubular extension 607 fitting within the lower portion of
spout 605. A spring 608 is under compression and is held
35 in position by the upper conical surface 609 of piston 606

1 and an inner shelf 610 of the spout 605. Lower pistons
611 and 612 are positioned within the lower portions of
compartments 602 and 603 respectively so as to follow the
dental material upwardly as it is being dispensed into the
5 spouts 604 and 605 and eventually into nozzle 613. The
upper part of container 601 has a reduced diameter
to encircle the nozzle 613 and provide for a sliding
engagement. Outer cap member 614 is threadedly engaged as
at 615, with outer surface of nozzle 613 to effectively
10 seal the container and prevent inadvertent dispensing of
dental material as well as a pin 616 which fits snugly
into open end of nozzle 613.

In operation, and with cap 614 removed, the user will
depress a push button lever 617 (seen in Fig. 7) located
15 outside the container 601. Lever 617 has substantially
flat elongated member 618 which projects between spouts
604 and 605 and presses against wall 619 which bridges
both spouts. Lever 617 is pivoted about pivot pin 620
affixed to inner wall of container 601. As lever 617
20 is depressed, member 618 will force spouts 604 and 605
downwardly and subsequently spring 608 as well as piston
606 will be lowered to phantom position (as seen in Fig.
6) causing dental material to flow upwardly within exten-
sion 607 and spout 605, mixing with material in spout 604
and through nozzle 613 to the bristles of a toothbrush.
25 As the lever is released, the spring 608 will force
nozzle 605 upwardly to its original position against
conical portion 621 of container 601. The vacuum created
will cause piston 606 to raise upwardly and concomitantly
cause lower piston 612 to travel upwardly the distance of
30 the expelled dental material. A spring clip 621 slideably
engages inner surface of compartment 603 to allow piston
to travel upwardly but be prevented from movement down-
wardly. Description of compartment 602 and spout 604 with
its accompanying component parts operate in a like manner
35 as described above.

The particular packaging arrangement used is not important. Those skilled in the art will be able to fashion several obvious modifications of the containers described herein by way of illustration.

Tubes, such as those suitable for use in accordance with the present invention are usually extruded around a cylindrical mandrel, cut into tube segments of suitable length, fitted with head structures and then filled from the bottom and pressed and/or welded closed, substantially as described in, e.g., U.S. Patent No. 4,060,179 issued on November 29, 1977 to McGhie, the disclosure of which is incorporated herein by reference.

In the case of the tube-within-a-tube embodiment of the present invention, the outer tube is provided first but it is not closed at the end opposite to that of the closure. The inner tube (also open-ended at the corresponding end) is inserted and fastened to the mouth of the outer tube. The two tubes are then filled and sealed together. A similar tube-within-a-tube arrangement has been proposed and described in U.S. Patent No. 1,566,218 of Leland and issued on December 15, 1925.

The invention is further illustrated by the following specific examples which are designed merely to illustrate the present invention and not to limit its scope.

In these examples, a hydrogen peroxide gel containing 3 - 6.5% hydrogen peroxide by weight, useful for simultaneous administration with a sodium bicarbonate paste is prepared as follows:

EXAMPLE 1

Ingredients

Hydrogen peroxide, 35% aqueous solution (5% H ₂ O ₂ in final gel)	14.3 parts
---	------------

35	Purified water	84.45
----	----------------	-------

- 1 Copolymer of acrylic acid crosslinked with
1% by weight of polyallyl sucrose having
5.8 allyl groups per molecule (CARBOPOL 934
made by B.F. Goodrich Chemical Co.,
5 Akron, Ohio) 0.5
Hydroxyethyl cellulose 0.5
Triethanolamine 0.25
- The gel is prepared by combining the hydrogen peroxide
solution with the purified water, followed by the gradual
10 addition of CARBOPOL 934. Upon thorough dispersion of the
copolymer hydroxyethyl cellulose is slowly added and
dissolved. Finally, triethanolamine is added, forming a
clear, homogeneous, stable and viscous gel having a pH of
3.4.
- 15 EXAMPLE 2
Ingredients
Hydrogen peroxide, 35% aq. solution (3.5% H_2O_2
in final gel) 10.0 parts
Distilled or deionized water 88.9
20 Acrylic acid copolymer CARBOPOL 940 (Goodrich) 0.6
Hydroxyethylcellulose 0.5
Sodium hydroxide, 10% solution q.s. pH 3.8 - 4.0
Preparation: same as that of Example 1.
- EXAMPLE 3
Ingredients
25 Hydrogen peroxide, 35% (3.5% H_2O_2 in final gel) 10.0 parts
Distilled or deionized water 89.0
Acrylic acid copolymer - CARBOPOL 941 (Goodrich) 0.7
Hydroxypropylcellulose 0.3
30 Sodium hydroxide, 10% solution q.s. pH 3.8 - 4.0
Preparation: same as that of Example 1.
- EXAMPLE 4
Ingredients
Hydrogen peroxide, 35% (4.0% H_2O_2
35 in final gel) 11.5 parts

-15-

1	Distilled or deionized water	86.65
	Acrylic acid copolymer - CARBOPOL 934 (Goodrich)	0.75
	Sodium laurylsulfate, dentifrice grade	0.50
	Hydroxypropylcellulose	0.6
5	Sodium hydroxide, 10% solution	q.s. pH 3.5 - 4.5

Preparation:

The hydrogen peroxide solution is combined with the distilled or deionized water. Sodium laurylsulfate is added under constant agitation and dissolved. Gradually, CARBOPOL 934 is added and dispersed. Hydroxypropylcellulose is added in increments and dissolved. When the mixture is homogeneous, sodium hydroxide is added slowly to the desired pH level and viscosity.

EXAMPLE 515 Ingredients

	Hydrogen peroxide, 35% (6.0% H_2O_2 in final gel)	17.14 parts
	Distilled or deionized water	81.76
	Acrylic acid copolymer - CARBOPOL 940 (Goodrich)	0.70
20	Hydroxyethylcellulose	0.40
	Sodium hydroxide, 10% solution	q.s. pH 3.5-4.0

Preparation: same as that of Example 1.

EXAMPLE 6Ingredients

25	Hydrogen peroxide, 35% (3.0% H_2O_2 in final gel)	8.58 parts
	Distilled or deionized water	89.22
	Acrylic acid copolymer - CARBOPOL 934 (Goodrich)	0.70
30	Hydroxypropyl methylcellulose	0.65
	Nonionic surfactant PLURONIC F 127 (BASF Corp., New Jersey)	0.85
	Sodium hydroxide, 10% solution	q.s. pH 3.5 - 4.5

Preparation: same as that of Example 4.

1 EXAMPLE 7

The sodium bicarbonate paste is prepared as follows:

Ingredients

	Deionized water	31.94 parts
5	Sorbitol 70% solution, USP	20.0
	Cellulose gum - CMC 7MF (Hercules)	1.44
	Sodium saccharin	0.20
	Magnesium aluminum silicate - VEEGUM F (made by R. T. Vanderbilt Co.,	
10	Inc., Norwalk, Conn.)	1.17
	Sodium bicarbonate, fine powder	40.00
	Sodium chloride	40.00
	Sodium lauryl sulfate - dentifrice grade	0.30
	Peppermint/Spearmint Flavor	0.75
15	Methyl paraben, USP	0.15
	Propyl paraben, USP	0.05

Procedure:

Glycerin and propylene glycol are combined in a first
20 container with agitation. Cellulose gum is added and
dispersed thoroughly throughout the mixture. Saccharin,
methylparaben and propylparaben are added in a separate
container and heated to dissolve. VEEGUM is added and the
mixture is agitated until uniform. The contents of the
25 first container are slowly added to the second container
and the final mixture is agitated thoroughly until
uniform. Flavoring agent, sodium lauryl sulfate and
coloring (if desired) are added and the paste is agitated
at moderate speed until uniform. Entrapped air is removed
30 by degassing in a vacuum vessel. Further homogeneity may
be obtained by milling, if necessary.

EXAMPLE 8Ingredients

	Deionized water	33.43 parts
35	Glycerin	10.00

- 1 Glycerin and propylene glycol are combined in a first
 container with agitation. Cellulose gum is added and
 dispersed thoroughly throughout the mixture. Saccharin,
 methylparaben and propylparaben are added in a separate
 5 container and heated to dissolve. VEEGUM is added and the
 mixture is agitated until uniform. The contents of the
 first container are slowly added to the second container
 and the final mixture is agitated thoroughly until
 uniform. Flavoring agent, sodium lauryl sulfate and
 10 coloring (if desired) are added and the paste is agitated
 at moderate speed until uniform. Entrapped air is removed
 by degassing in a vacuum vessel. Further homogeneity may
 be obtained by milling, if necessary.

EXAMPLE 815 Ingredients

	Deionized water	33.43 parts
	Glycerin	10.00
	Propylene glycol	10.0
	Cellulose gum - CMC 7MF (Hercules)	1.45
	Sodium saccharin	0.20
20	Magnesium aluminum silicate - VEEGUM F	1.17
	Sodium bicarbonate, fine powder	25.00
	Dicalcium phosphate dihydrate	13.50
	Dicalcium phosphate, anhydrous	1.50
	Sodium chloride	2.50
25	Sodium lauryl sulfate, dentifrice grade	0.30
	Methylparaben, USP	0.15
	Propylparaben, USP	0.05
	Peppermint/Spearmint Flavor	0.75
	FD & C Blue No. 1, 0.1% solution	q.s.
30	DS & C Yellow No. 6, 0.1% solution	q.s.

Procedure: Same as that of Example 8.

EXAMPLE 9

Paste containing fluoride:

35 Ingredients

	Deionized water	33.51
--	-----------------	-------

1	Sorbitol, 70% solution	20.00
	Sodium saccharin	0.20
	Cellulose gum CMC 7MF (Hercules)	1.54
	Magnesium aluminum silicate - VEEGUM F	1.17
5	Sodium fluoride	0.33
	Methyl paraben, USP	0.15
	Propyl paraben, USP	0.05
	Calcium sulfate	10.00
	Sodium bicarbonate	25.00
10	Sodium chloride	2.00
	Hydrated aluminum oxide	5.00
	Peppermint/Spearmint Flavor	0.75
	Sodium lauryl sulfate	0.30

Procedure: Same as that of Example 8.

15 EXAMPLE 10

Peroxide Gels

Composition 10-A

	Hydrogen peroxide, 35% aqueous solution (4.0% H_2O_2 in final gel)	11.5
20	Distilled water	86.6
	Acrylic acid copolymer - Carbopol 934 (Goodrich)	1.5
	Sodium lauryl sulfate, dentifrice grade	0.1
	Hydroxypropyl cellulose	0.3
	Sodium hydroxide, 10% solution q.s. pH 3.0 - 4.5	

25 Composition 10-B

	Hydrogen peroxide, 35% (4.0% H_2O_2 in final gel)	11.5
	Distilled water	88.0
	Acrylic acid copolymer (Carbopol 934, 940, 941, or 1342)	1.5
30	Sodium hydroxide, 10% solution q.s. pH 3.0 - 4.5	

Composition 10-C

	Hydrogen peroxide, 35% (4.0% H_2O_2 in final gel)	11.5
	Distilled water	46.0

1	Glycerin, anhydrous	40.0
	Acrylic acid copolymer - Carbopol 941	2.5

Methods of preparation

Composition 10-A: same as listed in Example 4

- 5 Composition 10-B: same as in Example 4 except that sod. lauryl sulfate and hydroxypropyl cellulose were omitted. The composition including Carbopol 1342 has not been actually made.

- 10 Composition 10-C: The glycerin and water were combined and heated to 50 - 60°C. Very slowly, Carbopol 941 was added under constant agitation.

- 15 When a clear gel had formed and no undissolved lumps remained, the gel was cooled to 25°C and the hydrogen peroxide was added. Agitation was maintained until the mixture became homogeneous. The gel was de-aerated in a vacuum vessel.

EXAMPLE 11: Sodium bicarbonate Paste

Composition 11-A

	Glycerin	25.0
20	Cellulose gum CMC 7MF (Hercules)	1.54
	Deionized water	32.71
	Magnesium aluminum silicate Veegum (R.T. Vanderbilt)	1.10
	Sodium saccharin	0.60
25	Sodium chloride	2.0
	Methyl paraben	0.15
	Propyl paraben	0.05

(Sodium hydroxide solution 10%, q.s. pH 8.0 - 8.5, may be added, if necessary for pH adjustment)

30	Sodium fluoride	0.22
	Bentonite	4.0
	Titanium dioxide	2.0
	Silica	4.0
	Sodium bicarbonate	25.0

-20-

1 Flavor (spearmint) 1.0
Sodium lauryl sulfate 0.3
Color (FDC Blue No. 1) q.s.

Method of preparation

5 The cellulose gum was added to the glycerin and dispersed thoroughly.

In a separate vessel, the parabens, sodium saccharin, and sodium chloride were dissolved at 60 - 70°C. To the clear solution was added the Veegum and the mixture was agitated until uniform. The pH of this solution was determined to be 8.0 - 8.5 (and adjusted, if necessary).

The gum dispersion was added to the Veegum solution and agitated until uniform.

15 To the blend were added the powders, bentonite, TiO_2 , silica, NaHCO_3 , and NaF, under vigorous agitation.

To the paste were added flavor, sodium lauryl sulfate and color.

20 The finished paste is milled and degassed in a vacuum vessel.

EXAMPLE 13

Urea peroxide gel

Composition 13-A

25 Urea peroxide (35% H_2O_2 equivalent) 10.0 parts by weight
Glycerin, anhydrous 90.0

Method: Urea peroxide is slowly added under agitation to the anhydrous glycerin until a clear gel is formed which has a suitable consistency for filling into collapsible tubes.

Composition 13-B

Urea peroxide 10.0
Acrylic acid copolymers - Carbopol 941 1.5
Glycerin, anhydrous 88.5

35 The method is the same as that for 13-A except the urea peroxide and Carbopol 941 are both added to the glycerin solution.

1 Although the present invention has been
described with reference to preferred embodiments, those
of ordinary skill in the art will readily appreciate that
many additions, deletions, modifications and substitutions
5 are possible within the spirit of the present invention
and the scope of the following claims.

10

15

20

25

30

35

-22-

What is claimed is:

1. A composition useful in combatting gum disease comprising:

(a) a gel component comprising (i) about 0.1 - 10% by weight of hydrogen peroxide; (ii) about 0.05 - 5.0% by weight of a water-dispersible copolymer of acrylic acid cross-linked with polyallyl sucrose; (iii) about zero to about 2.0% by weight of a nonionic cellulose stabilizer; (iv) a neutralizing agent selected from the group consisting of sodium hydroxide, potassium hydroxide, triethanolamine, diisopropylamine and ammonia in an amount sufficient to raise the gel pH to about 3 - 6; and (v) purified water;

(b) a paste component comprising: (i) about 2-60% sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, $MgCl_2$, $MgSO_4$, Na_2SO_4 , and K_2SO_4 ; (iii) about 2-60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol and mixtures thereof; (iv) about 0.1 - 5% of a thickener stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium oxide and mixtures thereof; and (vi) purified water; said paste component and gel component being combined immediately prior to use.

2. The composition of claim 1, said paste component also comprising a fluorine-containing compound selected from the group consisting of NaF, KF, sodium monofluorophosphate, potassium monofluorophosphate, sodium fluorosilicate, sodium fluorozeonate and mixtures thereof in an amount sufficient to yield about 200 -3000 ppm of fluorine; said composition also being effective against caries.

-23-

3. The composition of claim 2, said paste further comprising about 0.1 - 2.5% of a foaming agent selected from the group consisting of sodium lauryl sulfate, sodium lauroyl sarcosinate, sodium coconut monoglyceride sulfonate, sodium N-methyl-N-palmitoyl lauride, a polysorbate, a poloxamer and mixtures thereof.

4. The composition of claim 3, said paste further comprising about 1 - 30% of an additional cleansing agent selected from the group consisting of calcium sulfate, calcium phosphate, hydrated aluminum oxide, calcium carbonate, magnesium carbonate, magnesium silicate and mixtures thereof.

5. The composition of claim 4 wherein said paste component also comprises a preservative, a coloring agent and a flavoring agent.

6. The composition of claim 1, wherein said gel component comprises about 3.0 - 6.5% hydrogen peroxide; about 1 - 3% of said copolymer; and about 0.3 - 1.5% of said neutralizing agent and wherein said paste component comprises about 20-40% of said sodium bicarbonate; about 2 - 4% of said salt; about 15 - 25% of said humectant; about 1.0 - 2.0% of said stabilizer; and about 1.5 - 20% of said stabilizing polishing agent.

7. A composition useful in combatting gum disease comprising:

(a) a non-neutralized gel component consisting essentially of about 0.1 - 10.0% of hydrogen peroxide; about 0.05 - 5.0% of acrylic acid copolymer cross-linked with polyallyl sucrose; about 2 - 80% of a polyol selected from the group consisting of glycerin, sorbitol (70% solution), polypropylene glycol, polyethylene glycol, an ethoxylated lower

-24-

fatty alcohol, a propoxylated lower fatty alcohol and mixtures thereof; and purified water;

(b) a paste component comprising: (i) about 2-60% sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, $MgCl_2$, $MgSO_4$, Na_2SO_4 , and K_2SO_4 ; (iii) about 2 - 60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol and mixtures thereof; (iv) about 0.1 - 5% of a thickener stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium oxide and mixtures thereof; and (vi) purified water; said paste component being combined with said gel component immediately prior to use.

8. The composition of claim 7, said paste component also comprising a fluorine-containing compound selected from the group consisting of NaF, KF, sodium monofluorophosphate, potassium monofluorophosphate, sodium fluorosilicate, sodium fluorozeirconate and mixtures thereof, in an amount sufficient to yield about 200 - 3000 ppm of fluorine.

9. The composition of claim 8, said paste further comprising about 0.1 - 2.5% of a foaming agent selected from the group consisting of sodium lauryl sulfate, sodium lauroyl sarcosinate, sodium coconut monoglyceride sulfonate, sodium N-methyl-N-palmitoyl lauride, a polysorbate, a poloxamer and mixtures thereof.

10. The composition of claim 9, said paste further comprising about 1 - 30% of an additional cleansing agent selected from the group consisting of calcium sulfate,

-25-

calcium phosphate, hydrated aluminum oxide, calcium carbonate, magnesium carbonate, magnesium silicate and mixtures thereof.

11. The composition of claim 10 wherein said paste component also comprises a preservative, a coloring agent and a flavoring agent.

12. The composition of claim 7, wherein said gel component contains about 3.0 - 6.5% hydrogen peroxide; and about 20 - 60% of said polyol and wherein said paste component comprises about 20-40% of said sodium bicarbonate; about 2 - 4% of said salt; about 15 - 25% of said humectant; about 1.0 - 2.0% of said stabilizer; and about 1.5 - 20% of said stabilizing polishing agent.

13. A composition useful for combatting gum disease comprising:

(a) a gel component comprising: (i) about 2 - 25% urea peroxide; (ii) about zero - 5.0% of an acrylic acid copolymer crosslinked with polyallyl sucrose; the balance being glycerin; and

(b) a paste component comprising: (i) about 2-60% sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, $MgCl_2$, $MgSO_4$, Na_2SO_4 , and K_2SO_4 ; (iii) about 2 - 60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol and mixtures thereof; (iv) about 0.1 - 5% of a thickener stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium oxide and mixtures thereof; and (vi) purified water, said paste and said gel component being combined immediately prior to use.

-26-

14. The composition of claim 13, said paste component also comprising a fluorine-containing compound selected from the group consisting of NaF, KF, sodium monofluorophosphate, potassium monofluorophosphate, sodium fluorosilicate, sodium fluorozeirconate and mixtures thereof in an amount sufficient to yield about 200 - 3000 ppm of fluorine.

15. The composition of claim 14, said paste further comprising about 0.1 - 2.5% of a foaming agent selected from the group consisting of sodium lauryl sulfate, sodium lauroyl sarcosinate, sodium coconut monoglyceride sulfonate, sodium N-methyl-N-palmitoyl lauride, a polysorbate, a poloxamer and mixtures thereof.

16. The composition of claim 15, said paste further comprising about 1 - 30% of an additional cleansing agent selected from the group consisting of calcium sulfate, calcium phosphate, hydrated aluminum oxide, calcium carbonate, magnesium carbonate, magnesium silicate and mixtures thereof.

17. The composition of claim 16 wherein said paste component also comprises a preservative, a coloring agent and a flavoring agent.

18. The composition of claim 13 wherein said gel component contains about 8 - 12% of urea peroxide; about 1-3% of said acrylic acid copolymer; and glycerin and wherein said paste component comprises about 20-40% of said sodium bicarbonate; about 2 - 4% of said salt; about 15 - 25% of said humectant; about 1.0 - 2.0% of said stabilizer; and about 1.5 - 20% of said stabilizing polishing agent.

19. The composition of claim 5, wherein said gel copolymer is selected from the group consisting of Carbopol 934, 940, 941 and 1314, said gel stabilizer is hydroxypropyl

-27-

cellulose and said neutralizer is NaOH; said salt is NaCl, said paste humectant is glycerin, said stabilizing polishing agent is selected from the group consisting of TiO_2 , silica, bentonite and mixtures thereof; said fluorinated compound is NaF, said foaming agent is sodium lauryl sulfate, and said preservative is selected from the group consisting of methyl and propyl paraben.

20. The composition of claim 11 wherein said gel copolymer is Carbopol 941, and polyol is glycerin; said salt is NaCl, said paste humectant is glycerin, said stabilizing polishing agent is selected from the group consisting of TiO_2 , silica, bentonite and mixtures thereof; said fluorinated compound is NaF, said foaming agent is sodium lauryl sulfate, and said preservative is selected from the group consisting of methyl and propyl paraben.

21. The composition of claim 17 wherein said polymer is Carbopol 941, said salt is NaCl, said paste humectant is glycerin, said stabilizing polishing agent is selected from the group consisting of TiO_2 , silica, bentonite and mixtures thereof; said fluorinated compound is NaF, said foaming agent is sodium lauryl sulfate, and said preservative is selected from the group consisting of methyl and propyl paraben.

22. A combination of a two-compartment container and a composition consisting of a gel component and a paste component and contained in said container, said combination being suitable for use in combatting gum disease, said container comprising:

(a) a first compartment containing said gel, said gel comprising (i) about 1 - 10% of hydrogen peroxide, (ii) about 0.05 - 5.0% of a water dispersible copolymer of acrylic acid crosslinked with polyallyl sucrose; (iii) about zero - 2.0% of a nonionic cellulose stabilizer; (iv) an amount

-28-

of a neutralizing agent sufficient to adjust the gel pH to about 3.0 - 6.0, said neutralizing agent being selected from the group consisting of NaOH, KOH, triethanolamine, diisopropylamine and ammonia; and (v) water; such that the gel liquifies immediately upon contact with a mildly alkaline environment containing a strong electrolyte, thereby causing the release of bactericidally effective amounts of nascent oxygen; said gel having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from said container; said first compartment having an orifice for dispensing controlled amounts of said gel;

(b) a second compartment containing said paste, said paste comprising (i) about 2 - 60% by weight of sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, $MgCl_2$, $MgSO_4$, Na_2SO_4 , K_2SO_4 and mixtures thereof; (iii) about 2 - 60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol, and mixtures thereof; (iv) about 0.1 - 5.0% of a thickener-stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium dioxide and mixtures thereof; and (vi) purified water; said paste having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from said container, said second compartment having an orifice for dispensing controlled amounts of said paste substantially simultaneously with the dispensation of said gel; said first compartment orifice and said second compartment orifice being adapted to dispense said gel and said paste respectively at the same use point, said first and second compartments having a common wall portion and said orifices being substantially adjacent.

-29-

23. A combination of a two compartment container and a composition consisting of a gel component and a paste component, said composition being contained in said container, said combination being suitable for combatting gum disease, said container comprising:

(a) a first compartment containing said gel, said gel being a non-neutralized gel and comprising about 0.1 - 10% of hydrogen peroxide; about 0.05 - 5.0% of an acrylic acid copolymer crosslinked with polyallyl sucrose; about 2 - 80% of a polyol selected from the group consisting of glycerin, sorbitol 0 - 70% solution, polypropylene glycol, polyethylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol and mixtures thereof; and purified water; such that the gel liquifies immediately upon contact with a mildly alkaline environment containing a strong electrolyte, thereby causing the release of bactericidally effective amounts of nascent oxygen; said gel having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from said container; said first compartment having an orifice for dispensing controlled amounts of said gel;

(b) a second compartment containing said paste, said paste comprising (i) about 2 - 60% by weight of sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, $MgCl_2$, $MgSO_4$, Na_2SO_4 , K_2SO_4 and mixtures thereof; (iii) about 2 - 60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol, and mixtures thereof; (iv) about 0.1 - 5.0% of a thickener-stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium dioxide and mixtures thereof; and (vi) purified water; said paste having

-30-

sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from said container, said second compartment having an orifice for dispensing controlled amounts of said paste substantially simultaneously with the dispensation of said gel; said first compartment orifice and said second compartment orifice being adapted to dispense said gel and said paste respectively at the same use point, said first and second compartments having a common wall portion and said orifices being substantially adjacent.

24. A combination of a two compartment container and a composition consisting of a gel component and a paste component, said composition being contained in said container, said combination being suitable for combatting gum disease, said container comprising:

(a) a first compartment containing said gel, said gel comprising about 2 - 25% urea peroxide, about zero - 3.5% of an acrylic acid copolymer crosslinked with polyallyl sucrose; and glycerin; such that the gel liquifies immediately upon contact with a mildly alkaline environment containing a strong electrolyte, thereby causing the release of bactericidally effective amounts of nascent oxygen; said gel having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from said container; said first compartment having an orifice for dispensing controlled amounts of said gel;

(b) a second compartment containing said paste, said paste comprising (i) about 2 - 60% by weight of sodium bicarbonate; (ii) about 0 - 6% of a salt selected from the group consisting of NaCl, KCl, $MgCl_2$, $MgSO_4$, Na_2SO_4 , K_2SO_4 and mixtures thereof; (iii) about 2 - 60% of a humectant selected from the group consisting of glycerin, sorbitol, polyethylene glycol, polypropylene glycol, an ethoxylated lower fatty alcohol, a propoxylated lower fatty alcohol, and mixtures

-31-

thereof; (iv) about 0.1 - 5.0% of a thickener-stabilizer selected from the group consisting of cellulose gum, magnesium aluminum silicate and mixtures thereof; (v) about 1 - 30% of a stabilizing polishing agent selected from the group consisting of bentonite, titanium dioxide, silica, magnesium dioxide and mixtures thereof; and (vi) purified water; said paste having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from said container, said second compartment having an orifice for dispensing controlled amounts of said paste substantially simultaneously with the dispensation of said gel; said first compartment orifice and said second compartment orifice being adapted to dispense said gel and said paste respectively at the same use point, said first and second compartments having a common wall portion and said orifices being substantially adjacent.

25. The combination of claim 22, wherein said gel contains greater than 1.2 and up to about 5.0% of said copolymer and said two-compartment container is selected from the group consisting of two-compartment collapsible tubes with flexible sidewalls, two-compartment pressurized containers, two-compartment pumps, and combinations of two single-compartment tubes.

26. The combination of claim 23, wherein said two-compartment container is selected from the group consisting of two-compartment collapsible tubes with flexible sidewalls, two-compartment pressurized containers, two-compartment pumps, and combinations of two single-compartment tubes.

27. The combination of claim 24, wherein said two-compartment container is selected from the group consisting of two-compartment collapsible tubes with flexible sidewalls, two-compartment pressurized containers, two-compartment pumps, and combinations of two single-compartment tubes.

-32-

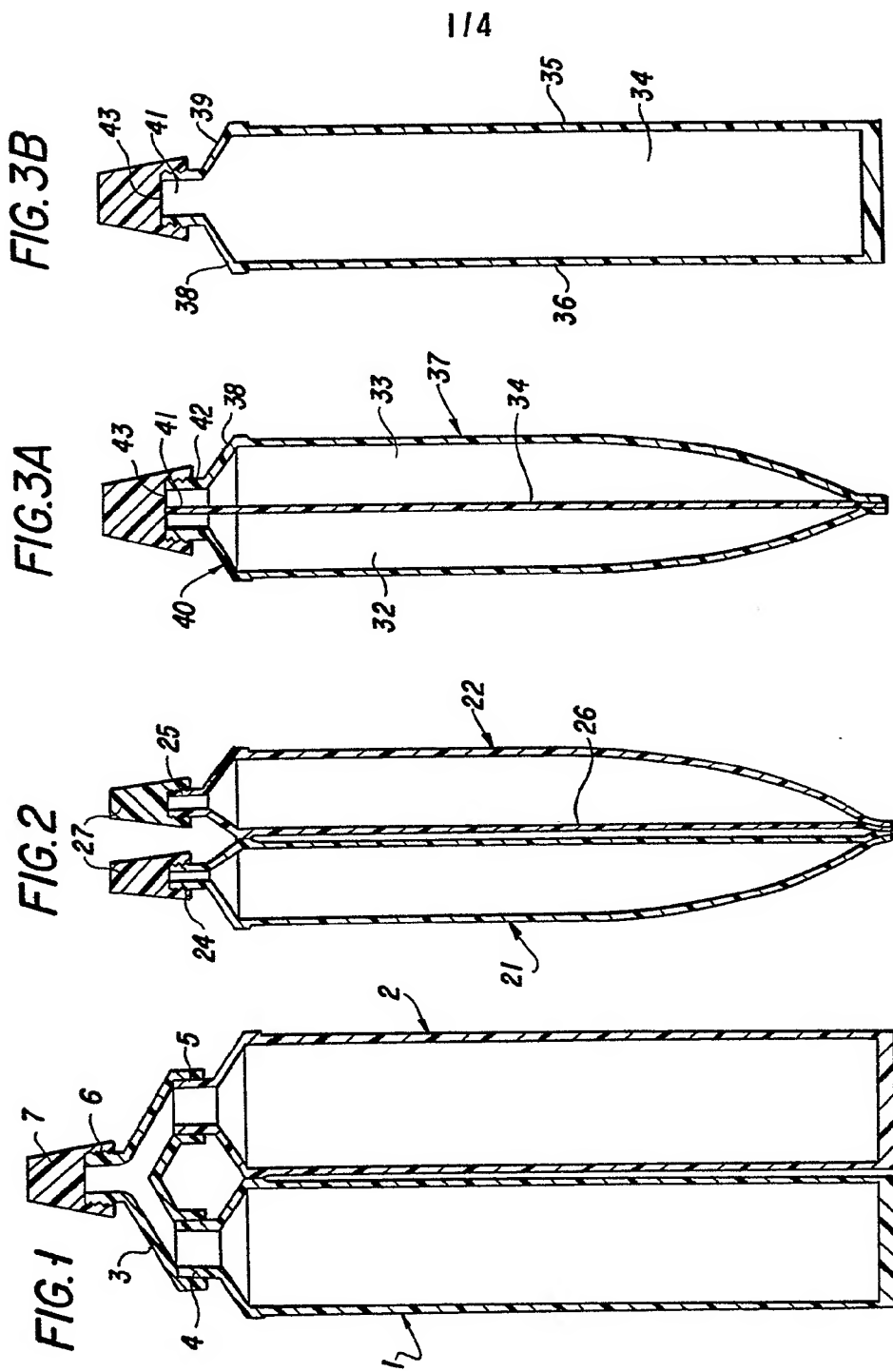
28. A composition useful in combating gum disease comprising:

(a) a gel component comprising an effective amount of peroxide selected from the group consisting of hydrogen peroxide and urea peroxide; a water dispersible polymer selected from the group consisting of a copolymer of acrylic acid cross-linked with polyallyl sucrose, a polyuronic acid colloid, a carboxymethylene colloid, a partially hydrolyzed polyacrylate, a partially hydrolyzed polymethacrylate, a partially hydrolyzed acrylate-methacrylate copolymer, and a polyoxyethylene-polyoxypropylene block copolymer; a nonionic cellulose gum stabilizer; purified water; and an amount of a neutralizing agent selected from the group consisting of: sodium hydroxide, potassium hydroxide, triethanolamine, diisopropylamine and ammonia sufficient to raise the pH of said gel to within about 3-6; said gel having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from a collapsible sidewall tube upon squeezing; and

(b) a paste component comprising effective amounts of sodium bicarbonate and a salt selected from the group consisting of sodium chloride and magnesium sulfate, at least one thickener-stabilizer selected from the group consisting of cellulose gum and magnesium-aluminum silicate, a bodying agent, purified water, at least one cleansing-polishing agent selected from the group consisting of calcium sulfate, calcium phosphate and hydrated aluminum oxide, and a foaming agent, said paste having sufficient viscosity to support itself on the bristles of a toothbrush and sufficient fluidity to be dispensed from a collapsible sidewall tube by squeezing, said gel and said paste components being mixed immediately prior to use.

-33-

29. The composition of claim 28 wherein said gel component comprises about 1-10% H_2O_2 , 0.05 - 1.2% acrylic acid copolymer and 0.1 - 1.5% non-ionic cellulose stabilizer by weight, said cellulose stabilizer being selected from the group consisting of hydroxyethyl cellulose, hydroxypropyl cellulose and hydroxypropyl methylcellulose, and wherein said paste component comprises about 10-50% sodium bicarbonate, 0-6% salt, 1-3% of at least one of cellulose gum and magnesium-aluminum silicate, 5-30% of a bodying agent, 1-40% of said cleanser-polishing agent and 0.1-2.5% foaming agent by weight.



2/4

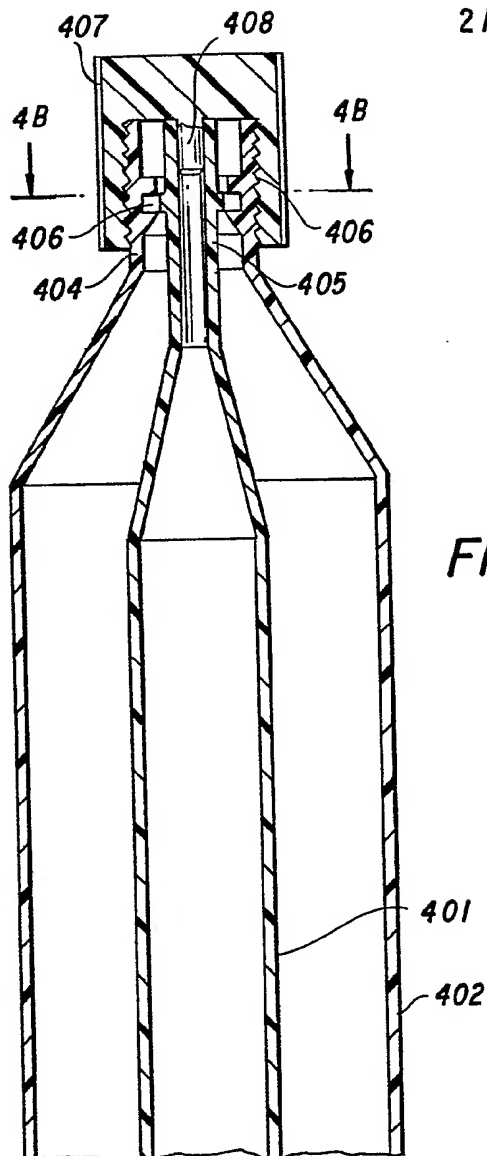
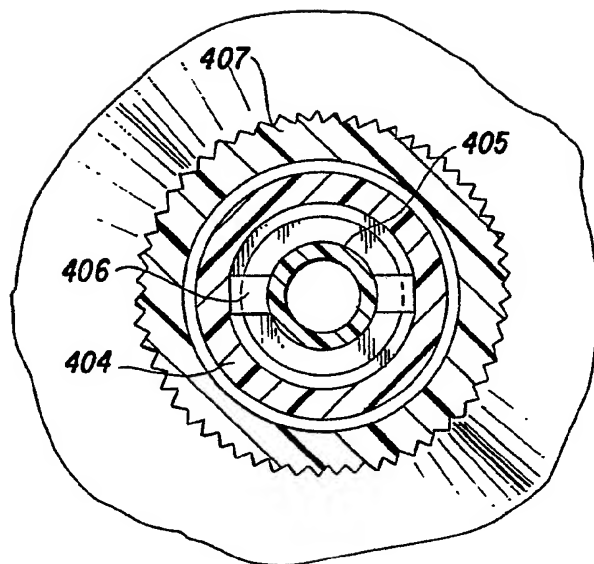
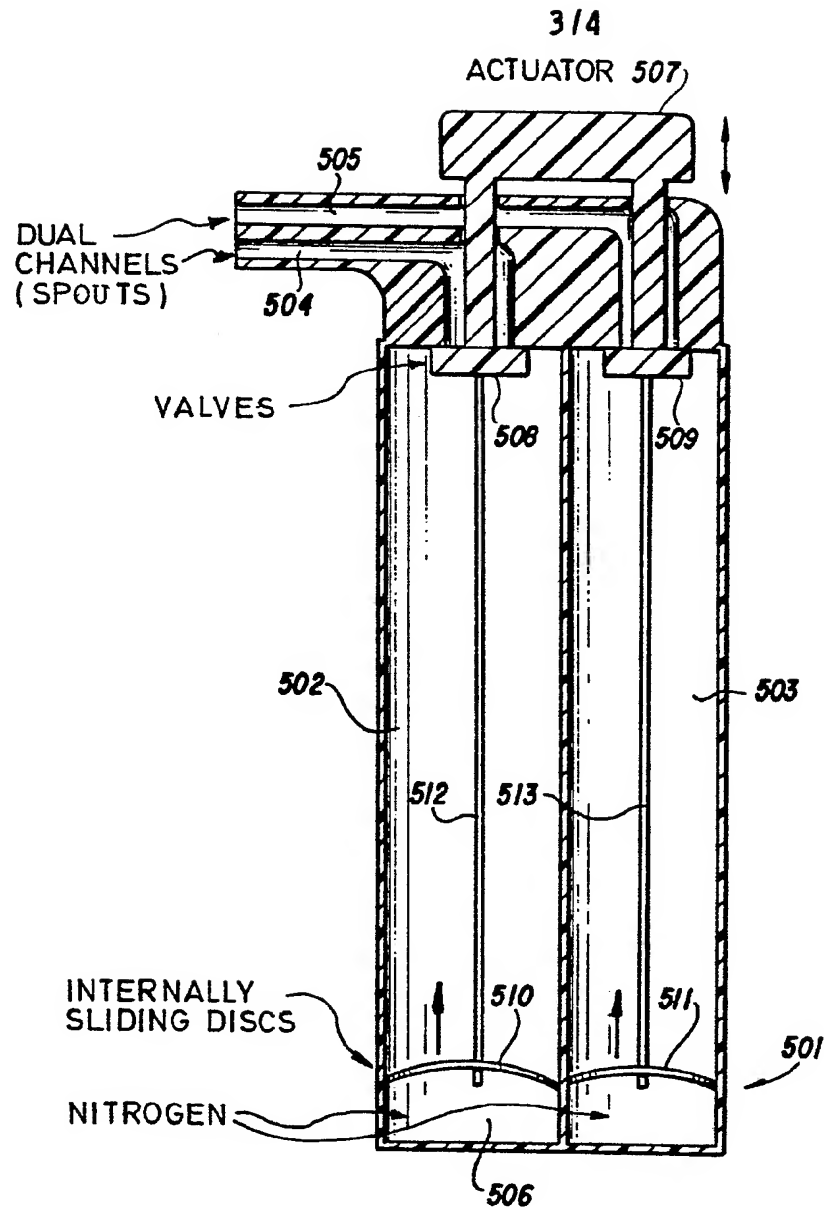


FIG. 4B



**FIG. 5**

DOUBLE COMPARTMENT PRESSURIZED CONTAINER

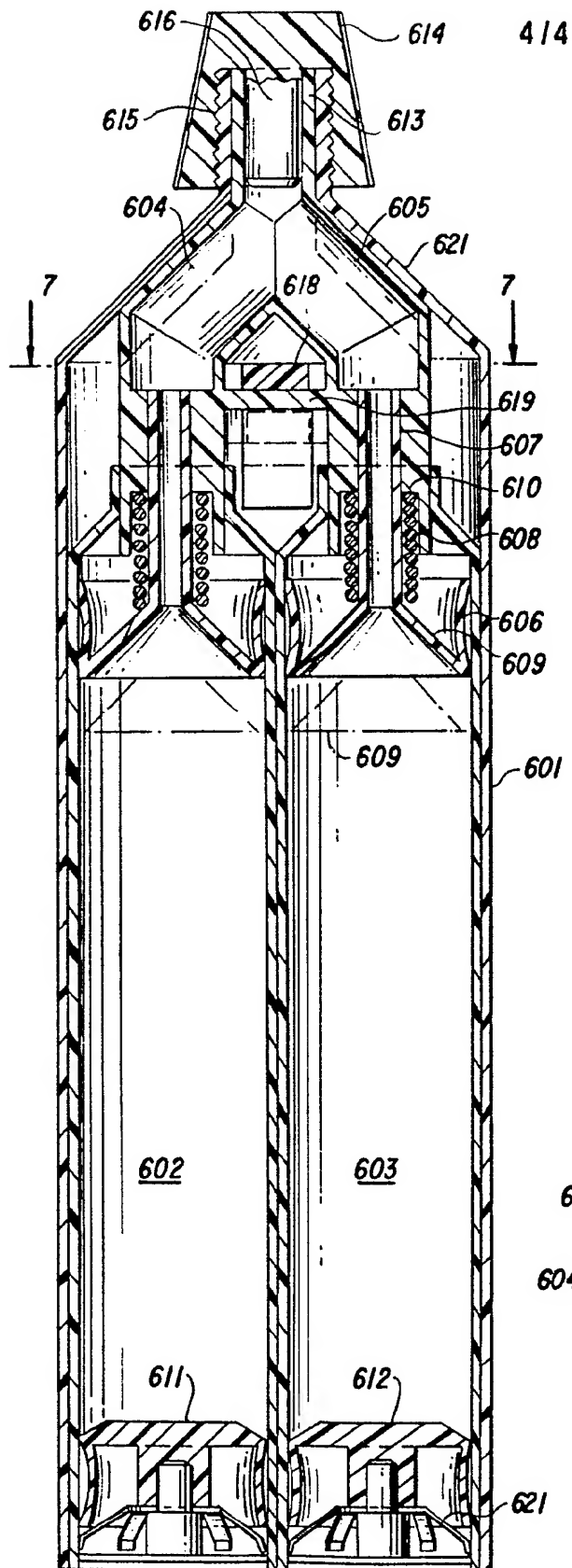


FIG. 6

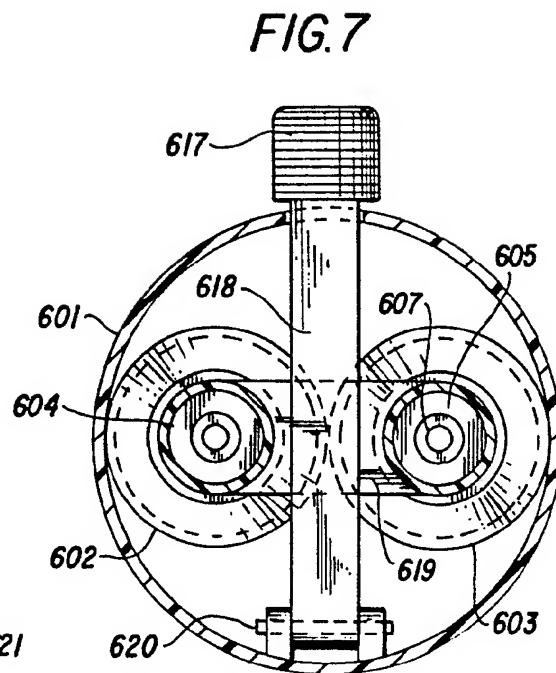


FIG. 7